## IN THE CLAIMS:

- (currently amended) A plant protein fraction which is derived from representatives of the Papaveraceae family and which possesses phospholipase D activity, comprising characterized in that
  - a) it consists of two protein subfractions A and B; B, and
  - b) it can be activated by Zn<sup>2+</sup> ions, and also
  - c) at least one of the subfractions A or and/or B possess carbohydrate wherein only protein subfraction A possesses moieties,

and with the protein subfraction A only possessing hydrolysis activity.

- (currently amended) The protein fraction as claimed in claim 1, characterized in that it
  is derived from Papaver somniferum and very particularly preferably from developing
  seedlings or and/or endosperms.
- 3 (currently amended) The protein fraction as claimed in claim 1 wherein or 2, eharacterized in that the subfraction A possesses a molecular mass of between 116 and 118 kDa, an isoelectric point, pI, of between 8.5 and 8.9 and a hydrolytic activity optimum at pH values of between 7.8 and 8.2, and the subfraction B possesses a molecular mass of between 112 and 115 kDa, an isoelectric point, pI, of between 6.5 and 6.9 and a hydrolytic activity optimum at pH values of between 5.0 and 6.0.
- 4. (currently amended) The protein fraction as claimed in <u>claim 1 wherein any one</u> of claims 1 to 3, characterized in that the subfraction A has a molecular mass of 116.4 kDa, an isoelectric point, pI, of 8.7 and a hydrolytic activity optimum at pH 8.0.

- 5. (currently amended) The protein fraction as claimed in <u>claim 1</u>, <u>wherein any one</u> of claims 1 to 4, characterized in that the subfraction B has a molecular mass of 114.1 kDa, an isoelectric point, pI, of 6.7 and a hydrolytic activity optimum at pH 5.5.
- 6. (currently amended) The protein fraction as claimed in <u>claim 1</u>, wherein any one of claims 1 to 5, characterized in that the subfraction B possesses an activatability optimum at Zn<sup>2+</sup> ion concentrations of between 1.0 and 10 mM and, particularly preferably, at 5 mM.
- 7. (currently amended) The protein fraction as claimed in <u>claim 1</u>, <u>wherein</u> <del>any one of claims 1 to 6, characterized in that</del> the subfractions A and B are isoenzymes.
- 8. (currently amended) The protein fraction as claimed in <u>claim 1</u>, <u>wherein</u> any one of claims 1 to 7, characterized in that its transphosphatidylating activity of the protein fraction is more strongly pronounced than its hydrolysis activity.
- 9. (currently amended) A method comprising The use of the protein fraction as claimed in any one of claims 1 to 8 for hydrolyzing or and/or transphosphatidylating phospholipids or and/or their lyso forms with the protein fraction of claim 1.
- 10. (currently amended) The method use as claimed in claim 9 for synthesizing phosphatidylcholine, phosphatidylethanolamine, phosphatidylglycerol, phosphatidylinositol, phosphatidic acid and phosphatidylserine and their lyso forms.
- 11. (currently amended) The method use as claimed in claim 9 or 10 in the form of a hydrolysis of phosphatidylinositol is hydrolized or and/or a headgroup exchange is performed on phosphatidylinositol.